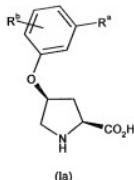


Amendments to the Claims:

1. - 10. (Canceled)

11. (Currently Amended) A pharmaceutical composition comprising a compound of formula (I)(la), as described in according to claim 114, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.
12. (Currently Amended) A combination comprising a compound of formula (I)(la), as described in according to claim 114, or a pharmaceutically acceptable salt, solvate or pre-drug thereof, and at least one other therapeutically active agent.
13. (Original) A combination according to claim 12, wherein the other therapeutically active agent is a PDEV inhibitor.

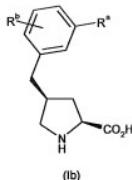
14. (New) A compound of formula (Ia):



wherein R<sup>a</sup> is selected from halogen, hydroxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, cyano, nitro, amino, hydroxycarbonyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkenyl, C<sub>1</sub>-C<sub>6</sub> alkynyl, hydroxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkoxyC<sub>1</sub>-C<sub>6</sub> alkyl, perfluoro C<sub>1</sub>-C<sub>6</sub> alkyl, perfluoroC<sub>1</sub>-C<sub>6</sub> alkoxy, C<sub>1</sub>-C<sub>6</sub> alkylamino, di- C<sub>1</sub>-C<sub>6</sub> alkylamino, aminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acyl, C<sub>1</sub>-C<sub>6</sub> acyloxy, C<sub>1</sub>-C<sub>6</sub> acyloxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acylamino, C<sub>1</sub>-C<sub>6</sub> alkylthiocarbonyl, C<sub>1</sub>-C<sub>6</sub> alkylthioxo, C<sub>1</sub>-C<sub>6</sub> alkoxy carbonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonylamino, aminosulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl;

R<sup>b</sup> is selected from hydrogen, halogen, hydroxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxy cyano, nitro, amino, hydroxycarbonyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkenyl, C<sub>1</sub>-C<sub>6</sub> alkynyl, hydroxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkoxyC<sub>1</sub>-C<sub>6</sub> alkyl, perfluoro C<sub>1</sub>-C<sub>6</sub> alkyl, perfluoroC<sub>1</sub>-C<sub>6</sub> alkoxy, C<sub>1</sub>-C<sub>6</sub> alkylamino, di- C<sub>1</sub>-C<sub>6</sub> alkylamino, aminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acyl, C<sub>1</sub>-C<sub>6</sub> acyloxy, C<sub>1</sub>-C<sub>6</sub> acyloxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acylamino, C<sub>1</sub>-C<sub>6</sub> alkylthio, C<sub>1</sub>-C<sub>6</sub> alkylthiocarbonyl, C<sub>1</sub>-C<sub>6</sub> alkylthioxo, C<sub>1</sub>-C<sub>6</sub> alkoxy carbonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonylamino, aminosulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

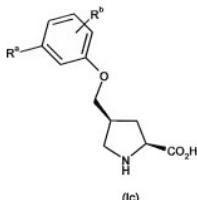
15. (New) A compound of formula (Ib):



wherein R<sup>a</sup> is selected from halogen, hydroxy, cyano, nitro, amino, hydroxycarbonyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkenyl, C<sub>1</sub>-C<sub>6</sub> alkynyl, hydroxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkoxyC<sub>1</sub>-C<sub>6</sub> alkyl, perfluoroC<sub>1</sub>-C<sub>6</sub> alkoxy, C<sub>1</sub>-C<sub>6</sub> alkylamino, di-C<sub>1</sub>-C<sub>6</sub> alkylamino, aminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acyl, C<sub>1</sub>-C<sub>6</sub> acyloxy, C<sub>1</sub>-C<sub>6</sub> acyloxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acylamino, C<sub>1</sub>-C<sub>6</sub> alkylthio, C<sub>1</sub>-C<sub>6</sub> alkylthiocarbonyl, C<sub>1</sub>-C<sub>6</sub> alkylthioxo, C<sub>1</sub>-C<sub>6</sub> alkoxy carbonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonylamino, aminosulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl;

R<sup>b</sup> is selected from hydrogen, halogen, hydroxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, cyano, nitro, amino, hydroxycarbonyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkenyl, C<sub>1</sub>-C<sub>6</sub> alkynyl, hydroxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkoxyC<sub>1</sub>-C<sub>6</sub> alkyl, perfluoro C<sub>1</sub>-C<sub>6</sub> alkyl, perfluoroC<sub>1</sub>-C<sub>6</sub> alkoxy, C<sub>1</sub>-C<sub>6</sub> alkylamino, di-C<sub>1</sub>-C<sub>6</sub> alkylamino, aminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acyl, C<sub>1</sub>-C<sub>6</sub> acyloxy, C<sub>1</sub>-C<sub>6</sub> acyloxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acylamino, C<sub>1</sub>-C<sub>6</sub> alkylthio, C<sub>1</sub>-C<sub>6</sub> alkylthiocarbonyl, C<sub>1</sub>-C<sub>6</sub> alkylthioxo, C<sub>1</sub>-C<sub>6</sub> alkoxy carbonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonylamino, aminosulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

16. (New) A compound of formula (Ic):



(Ic)

wherein R<sup>a</sup> and R<sup>b</sup> are independently selected from hydrogen, halogen, hydroxy, (C<sub>1</sub>-C<sub>6</sub>)alkoxy, cyano, nitro, amino, hydroxycarbonyl, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkenyl, C<sub>1</sub>-C<sub>6</sub> alkynyl, hydroxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkoxyC<sub>1</sub>-C<sub>6</sub> alkyl, perfluoro C<sub>1</sub>-C<sub>6</sub> alkyl, perfluoroC<sub>1</sub>-C<sub>6</sub> alkoxy, C<sub>1</sub>-C<sub>6</sub> alkylamino, di- C<sub>1</sub>-C<sub>6</sub> alkylamino, aminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminoC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acyl, C<sub>1</sub>-C<sub>6</sub> acyloxy, C<sub>1</sub>-C<sub>6</sub> acyloxyC<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> acylamino, C<sub>1</sub>-C<sub>6</sub> alkylthio, C<sub>1</sub>-C<sub>6</sub> alkylthiocarbonyl, C<sub>1</sub>-C<sub>6</sub> alkylthioxo, C<sub>1</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylsulfonylamino, aminosulfonyl, C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, di-C<sub>1</sub>-C<sub>6</sub> alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

17. (New) A compound of formula (Ia) according to claim 1 which is:

(2S, 4S)-4-(3-Chloro-phenoxy)-pyrrolidine-2-carboxylic acid;  
or a pharmaceutically acceptable salt thereof.

18. (New) A compound of formula (Ib) according to claim 15 which is selected from the group consisting of:

(2S,4S)-4-(3-Fluoro-benzyl)-pyrrolidine-2-carboxylic acid;  
(2S,4S)-4-(2,3-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; and  
(2S,4S)-4-(2,5-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; or a pharmaceutically acceptable salt thereof.

19. (New) A compound of formula (Ic) according to claim 16 which is selected from the group consisting of:

(2S,4S)-4-(3-Fluoro-phenoxyethyl)-pyrrolidine-2-carboxylic acid;

(2S,4S)-4-(3,6-Difluoro-phenoxyethyl)-pyrrolidine-2-carboxylic acid;  
(2S,4S)-4-(2,3-Difluoro-phenoxyethyl)-pyrrolidine-2-carboxylic acid; and  
(2S,4S)-4-(3-Methoxy-phenoxyethyl)-pyrrolidine-2-carboxylic acid; or a pharmaceutically acceptable salt thereof.

20. (New) A pharmaceutical composition comprising a compound of formula (Ib) according to claim 15, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.

21. (New) A combination comprising a compound of formula (Ib) according to claim 15, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.

22. (New) A combination according to claim 21, wherein the other therapeutically active agent is a PDEV inhibitor.

23. (New) A pharmaceutical composition comprising a compound of formula (Ic) according to claim 16, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.

24. (New) A combination comprising a compound of formula (Ic) according to claim 16, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.

25. (New) A combination according to claim 24, wherein the other therapeutically active agent is a PDEV inhibitor.

26. (New) The compound (2S,4S)-4-(3-Chloro-phenoxy)-pyrrolidine-2-carboxylic acid or a pharmaceutically acceptable salt thereof.